KISSIN, Mikhail Lakovich, dotsent, kandidat tekhnicheskikh nauk, [deceased];

MINULA.V., landmer, retsensent; UL'IANIMSKIY,S.V., professor, doktor
tekhnicheskikh nauk, retsensent; UVINTSKY,O.N., inzhener, retsensent,
redaktor; GOIUBENKOVA,L.A., redaktor; MEDVEDHY,L.Ya., tekhnicheskiy
redaktor

[Heating and ventilating] Otoplenie i ventiliateiia. Isd.2-oe, perer.
Moskva, Gos.isd-vo lit-ry po stroitel'stvu i arkhitekture. Pr.1.
[Heating] Otoplenie. 1955. 390 p. (MIRA 9:3)

(Heat engineering)

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

1. 行型的现在特殊型的翻譯· 网络克尔

KAMENEY, P.N., doktor tekhnicheskikh nauk, professor; GAMBURG, P.Tu., kandidet tekhnicheskikh nauk, dotsent; KISSIN, M.I., kandidet tekhnicheskikh nauk, dotsent [deceased]; SHCHEGLOV, V.P., kandidet tekhnicheskikh nauk, dotsent; STAROVEROV, I.G., inshener, retsensent; HIMMYAGI, D.K., redektor isdatel*stva; PERSON, M.N., tekhnicheskiy redaktor

[Heating and ventilation] Otoplenie i ventiliatsiia. Moskva, Gos. isd-vo lit-ry po stroit. i arkhit. Pt.l. [Heating] Otoplenie. 1956. 343 p. (MIRA 1012)

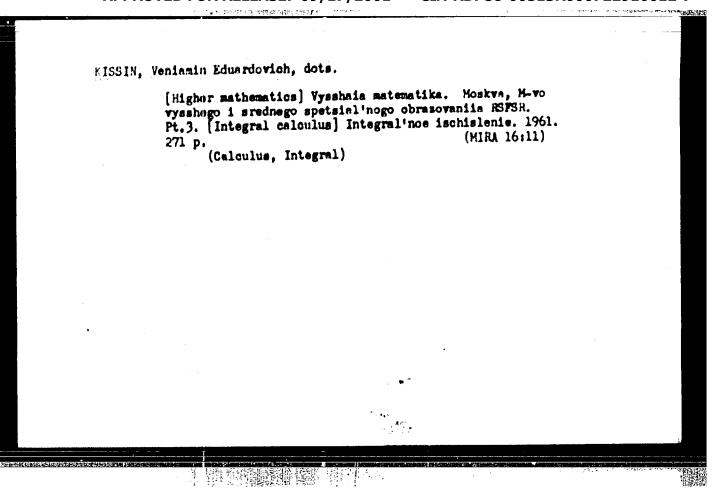
(Heat engineering)

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

- Contract the contract of the second of

KISSIE, M.L. kandidat tekhnicheskikh nauk, dotsent; D'YAKOHOV, P.I., kandidat tekhnicheskikh nauk, dotsent, retsensent; UL'YAKIESKIY, S.V., professor, retsensent; TURKUS, A.V., dotsent, redaktor; DAKHNOY, Y.S., tekhnicheskiy redaktor.

> [Heating and ventilation] Otoplenie i ventiliateiia. Pt. 1. [Heating] Otoplenie. Moskva, Gos. isd-vo stroit. lit-ry, 1947. 353 p. (MIRA 8:2) (Heating)



s/069/61/023/003/003/004 B127/B217

AUTHORS:

Khodzhayeva, I. V., Kissin, Yu. V.

TITLE:

Propresentation of the Control of th

Radiochromatographic separation of mixtures of sulfur and

vulcanization accelerators

PERIODICAL:

Kolloidnyy shurnal, v. 23, no. 3, 1961, 322-326

TEXT: The authors used the method of paper chromatography for the separation of radioactively tagged substances. It permits working with smallest quantities of the dangerous substances. Tetramethyl thiuramdisulfide (1) + S^{35} , the salt of diethyl dithiocarbamic acid (2) + S^{35} , and mixtures of tetraethyl thiuramdisulfide (TEDS) and (2) were studied. TEDS contained 4 atoms S^{35} (3). The reason for the selection was the frequent use of (1) and of some salts of (2) as vulcanization accelerators. It is assumed that the vulcanization activity is closely connected with the mobility of S atoms, in the molecules. The isotope exchange between (1) and S²⁵, (2) and S²⁵ took place by heating their solutions in benzene or in chloroform in sealed ampuls at 120-180°C. The exchange without a solvent was studied as well. Card 1/4

3/069/61/023/003/003/004 B127/B217

Radiochromatographic separation of ...

Methanol/H20/CH3COOH = 8:1:1 was used as a flux for (1), petroleum ether/ H20/CH3COOH = 8:1:1 for (2). The salts of Co, Ni, Cu, Cd, Pe, Pb, Hg of (2) were synthesized by precipitation with the respective cations from aqueous Na solution of (2) at certain pH. TETD* was produced by the method of Rothstein and Binovic (Recueil trav. chim. 73, 561, 1954) for the exchange between (2) and tagged thiurams. The reaction was carried out at 25°C in CHCl, at a TETD* concentration of 0.04 moles/1 and the molar ratios 1:1.5 for Co-+Fe salts of (2) to TETD*. When the ampuls containing the solutions of S35 and (2) were heated to more than 100°C, (2) was decomposed under sulfide precipitation. For the separation of the mixture, a small part of the solution (0.005-0.01 moles containing 10-30y of the substance) was dropped on a special paper filter strip of 40 cm length and 4.5 cm width. The activity of the spot was 2000-3000 decays/min. Then, the strip was dried and put together with the flux into the chromatographic chamber. Por evaluating the chromatogram, the paper strip was subdivided into 1-2 cm long sections, and the activity of the individual parts measured by a Geiger counter. The results are shown in a diagram. The distance from the

Card 2/4

S/069/61/023/003/003/004 B127/B217

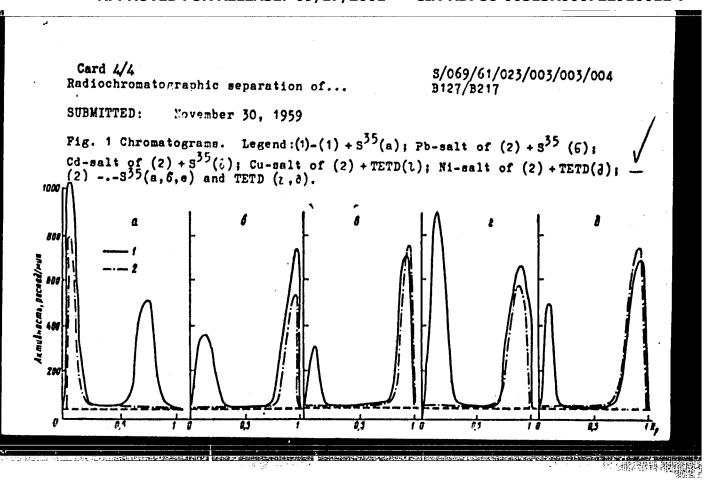
Radiochromatographic separation of ...

first spot was plotted on the x-axis, the activity on the y-axis (Fig.). The following formula was used for calculating the exchange percentage:

 $\%_{\text{exchange}} = 2\sum_{i=1}^{\infty} (I_{1} - I_{0}) / [\sum_{i=1}^{\infty} (I_{1} - I_{0}) + \sum_{i=2}^{\infty} (I_{2} - I_{0})].$ I_1 and I_2 are the maximum activities of the 1-2 cm long sections: I_0 is the activity on the background. Methyl- and ethyl alcohol as mobile phase, and water as steady phase were used as fluxes for the separation of (1) from 35, furthermore CH3COOH in order to increase the discrimination. Ethyl ether, H2O and CH3COOH (8:1:1) were the fluxes for the separation of TETD* from (2). The first activity is that of (2), the activity in the final spot is that of TETD*. The control experiment with pure TETD* in ethanol shows that the total amount of TETD* is concentrated in the final spot. There are 1 figure, 2 tables, and 5 references: 3 Soviet-bloc.

ASSOCIATION: Moskovskiy institut tonkoy khimicheskoy tekhnologii im. L. V. Lomonosova (Noscow Institute of Fine Chemical Technology imeni H. V. Lomonosov)

Chied 3/4



FIRSOV, A.P.; KASHPOROV, B.G.; KISSIN, Yu.V.; CHIRKOV, N.M.

Stereospecific action of the complex catalyst d_-TiOl3 - Me(C,Es)n in the polymorization of d_-olefins depending on the nature of the metal of the organometallic compound. Vymokom.soed. 4 no.7:1124
J1 '62. (Alra 15:7)

(Olefins) (Polymorization)

(Organometallic compounds)

8/020/62/145/001/013/018 B145/B101

Kissin, Yu. V., Tolstykh, E. V., and Chirkov, H. M.

Infrared spectra of the reaction products of (C5H5)2TiCl2 AUTHORS:

with aluminum alkyls TITLE:

PERIODICAL: Akademiya nauk SSSR. Doklady, v. 145, no. 1, 1962, 104 - 105 TEXT: The IR spectra of the "blue complexes" (C5H5)2TiCl2A1(C2H5)2 (I). (C5H5)2TiCl2Al(C2H5)Cl (II) and for comparison these of (C5H5)2TiCl2. Al(C2H5)3, and the dimer of Al(C2H5)2Cl were taken and are here discussed. The complexes were prepared by reaction of $(C_5H_5)_2$ TiCl with Al $(C_2H_5)_3$ $Al(C_2H_5)_2Cl$ in heptane. In the 1200 - 700 cm⁻¹, region the spectra of the complexes correspond to the sum of the spectra of (C5H5)2TiCl2 plus the corresponding aluminum alkyl. The intensive 870 cm band of (C5H5)2TiCl2 does not occur, whereas its 820 cm band is shifted to 812 - 810 cm and coincides with the absorption band of aluminum alkyl. The intent Card 1/2

CIA-RDP86-00513R000722910012

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                                                                                                                                                                                                                               s/190/63/005/001/009/020
                                                                                                                                                                                                                                 B101/B186
                                                                 Berdnikova, M. P., Kissin, Yu. V., Chirkov, N. M.
                                                                    Polymerization of a-amylene on complex catalysts
     PERIODICAL: Vysokomolekulysrnyye soyedineniya, v. 5, no. 1, 1963, 63-67
    5 27 5
        TEXT: The polymerization of 3-methyl-butene-1 and of n-pentene-1, both rick. The polymerization of 3-methyl-butene-1 and of n-pentene-1, both rick.
          TEXT: The polymerization of 3-methyl-butene-1 and of n-pentene-1, both ricl; catalyst is reported.

dissolved in n-heptane, with an Al(C2H5)3 70°C with a ratio of 70°C was at 40° 70°C with a ratio of 70°C was 3-methyl-butene-1 was polymerized at 40° 70°C was at 70°C was at 70°C was 3-methyl-butene-1 was polymerized at 40°C the reaction rate at 70°C was al (C2H2); TiCl. 1.7. The constant of the reaction rate at 70°C was all (C2H2); TiCl. 1.7.
AUTHOHS:
                 5-methyl-butene-1 was polymerized at 40 - 70°C with a ratio of 70°C was Al(C2H5)3 - 1.7. The constant of the reaction rate at 70°C was Al(C2H5)3 - 1.7. The constant of the reaction rate at 1.7.
  TITLE:
                   2.8.10-41/min.8 Ticl3. its temperature dependence followed the Arrhenius
                      equation, and the activation energy was 10 kcsl/mole. The polymer, a
                         white powder, m. p. 250 - 240°C, Oxidized intensively above bands insoluble in organic solvents, and did not form films.
                          white powder, m. p. 230 - 240°C, oxidized intensively above bands. The bands in organic solvents, and did not form films. 1460 cm band insoluble in its IR spectrum were the following: wibration of CH. groups and deformation vibration of CH.
                              identified in its IR spectrum were the following: the 1460 cm of CH<sub>2</sub> groups and deformation wibration of CH<sub>3</sub> groups and deformation of CH<sub>3</sub> groups are deformation of CH<sub>3</sub> groups and deformation of CH<sub>3</sub> groups are deformation of CH<sub>3</sub> groups and deformation of CH<sub>3</sub> groups are de
                                 asymmetric vibration of CH<sub>2</sub> groups and deformation vibration of CH<sub>2</sub> groups and deformation of CH<sub>3</sub> in the isopropyl a 1385 and 1385 card 1/3
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Polymerization of a-amylene on ...

S/190/63/U05/001/009/020 B101/B186

group. 1300 - 850 cm⁻¹ bands were not identified; they disappeared almost completely in the IR spectrum of the polymer melted at 260°C. They are perhaps caused by crystal interactions in the highly crystalline solid polymer. n-pentene-1 was polymerized at 70°C. The constant of the reaction rate was 2.3·10⁻³1/min·g TiCl₃. The polymer is a white, rubber-like and film-forming mass, m. p. 80°C; the shape of its deformation - stress curve is typical of elastomers. The following bands were identified in the IR spectrum: 1450 and 1370 cm⁻¹ bands as deformation vibrations of CH₃ and CH₂ groups, the 1340 cm⁻¹ band as deformation vibration of CH groups, the 1137 cm⁻¹ band as skeleton vibrations in branched polymer chains, the 1030 cm⁻¹ band as pendulum swings of CH₃ groups in the polymer side chains, the 1295 cm⁻¹ band as torsional vibrations of CH₂ groups, and the 727 cm⁻¹ band as pendulum swings of CH₂ groups. The 1640 cm⁻¹ band indicates the existence of double bonds in the end groups and the 956 cm⁻¹ band the existence of trans-double bonds. The formation of these Card 2/5

Polymerization of α -amylene on ...

\$/190/63/005/001/009/020 B101/B186

bands is explained by head-on-head addition besides head-on-tail addition of the monomer and termination in the resulting compound >Al-CH-(CH₂)₂-CH-R caused by steric hindrance owing to the removal of

one H atom from one methylene group of the principal or side chains. There

ASSOCIATION: Institut khimicheskoy fiziki AN SSSR .

(Institute of Chemical Physics AS USSR)

SUBMITTED:

July 17, 1961

Card 3/3

L 13543-63 EMP(1)/EPF(c)/EMT(m)/BDS ASD 9c-4/17-4 RM/WW ACCESSION NR: AP30C0685 8/0190/63/005/005/0653/0658 63-

AUTHOR: Pironoy, Q. M.; Kissin, Tu. V.; Chirkov, N. M.

14

TITLE: Synthesis and formation kinetics of low molecular poly-alpha-clefins on complex organometallic catalysts. 1. Polymerization of propylens in the presence of the catalytic system TiCl sub 4 and Al(iso-C sub 4 H sub 9)

SOURCE: Vy wookomolekulyarny wy soyedineniya, v. 5, no. 5, 1963, 633-638

TOPIC TAGS: synthesis, formation kinetics, poly-alpha-olefins, polymerization of propylene, catalytic systems

ABSTRACT: The present work was carried out to supply missing information on the polymerization kinetics of propylene over the systems Al(iso-C sub 4 H sub 9) sub 2 Cl and TiCl sub 4. Polymerization was conducted in high pressure installations at 34.8 to 17 atm and a temperature range of 60 to 100C, using liquid propane-propylene mixtures. Liquid polymers were obtained with a degree of polymerization ranging from 3 to 7 and higher. Their molecular weights depended on the temperature of polymerization and the Al:Ti ratio. Spectroscopic examination proved the polymers to be 100% olefins, with an approximate 5:1 ratio of the groups CH sub 2 = C (R) sub 2 and RHC = C(R) sub 2. A small amount of vinyl double bonds was also detected. The mechanism of double bond formation is discussed. Orig. art. has:

Cord 1/4 Association: Inst. of Chemical Physics, Academy of Sciences, SSSR

BM/WW Pc-li/Pr-li EMP(1)/EPF(c)/EMT(m)/BDS L 13553-63 8/0190/63/005/005/0719/0723 ACCESSION NR: AP3000700 AUTHOR: Romanov, L. M.; Verkhoturova, A. P.; Kissin, Tu. V.; Bakova, G.

TITLE: Polymerization of hepta-1,5-diene on complex catalysts

SOURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 5, no. 5, 1963, 719-723

ABSTRACT: The difficulties in obtaining rubbers suitable for vulcanization by means copolymerization of alpha-olefins induced the authors to select hepta-1,5-diene for a study of homopolymerization by means of various Ziegler-Hatta catalysts. The most active of these proved to be the system Al (C sub 2 H sub 5) sub 3 - TiCl sub 4. The Al/Ti ratio of 2:1 proved the most effective, producing a maximum 40% yield of the polymer in a n-heptime solution at 70 to 80C. The obtained poly-hepta-1,5-diene had a rubberlike texture, a molecular weight of 1250 and a 25-30% of double bonds, as determined by Harus' method. The product was also subjected to infrared spectroscopy in the 2000-7000 cm sup -1 range, and the number of double bonds per one CH sub 2 group was determined. Ozonization provided additional clues. It is concluded that the internal double bond is capable of participating in the polymerization initiated by Ziegler-Matta catalysts, a fact established by Matta while the present investigation was still in the experimental stage. The formation of five-membered Card 1/2

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Pc-4/Pr-4/Pt-4 RM/WW S/0190/63/005/007/1069/1071

ACCESSION NR: AP3(03796 S/0190/63/005/007/1069/1071

AUTHOR: Kissin, Yu. V.; Pshenitsy*na, G. M.

TITLE: Infrared spectra of polyaminoquinones

SOURCE: Vy*sokomolekulyarny*ye soyedineniya, v. 5, no. 7, 1963, 1069-1071

TOPIC TAGS: polymeric aminoquinone, polyaminoquinone, polyaminochloroquinone, benzidine, p-benzoquinone, chloranil, semiconductor, polymeric semiconductor, infrared spectroscopy, infrared spectra, conjugated bond system, bank shift, complex intramolecular complex, intermolecular complex

ABSTRACT: The structure of certain polymeric aminoquinones — reaction products of benzidine and r-benzoquinone or chloranil — has been investigated by infrared spectroscopy. The polymers were first synthesized by P. S. Shantarovich and G. M. Pshenitsyans (Vysokomolek. soyed., 5, no. 8, 1963), V. P. Parini et al. (Vysokomolek. soyed., 3, 402, 1961), and A. A. Berlin and Ye. G. Yatveyeva (Vysokomolek. soyed., 1, 1643, 1959) as potential polymeric semiconductors.) Shantarovich and Absorption spectra were measured in the 2000—700 cm⁻¹ region for KBr pellet samples. The reaction product of aniline and p-benzoquinone was used as a reference compound. As indicated by the spectra given in Fig. 1 of the Enclosure,

Card 1/4 2

L 12862-63 ACCESSION NR: AP30(13796

6

the band in the polymer of benzidine and p-benzoquinone due to c=0 is strongly shifted toward higher wavelengths with respect to the reference compound. This shift may be ascribed to the presence in the polymer chain either of quinoid-type groups or of groups containing disubstituted vinyl alcohol. The absence of a strong shift in the reference compound leads to the conclusion that in the polymer the intra- or intermolecular complexes responsible for the shift are stabilized by the conjugated-bond system. Polymers prepared with benzidine/p-benzoquinone ratios of 4/1 and 3/1 were both assigned the following structure:

The spectrum of the condensation product of benzidine and chloranil is in good agreement with the structure proposed by A. A. Berlin and Ye. G. Matkeyeva.

"The polymer samples were kindly made available to us by P. S. Shantarovich.
B. P. Parini, and N. G. Matyeyeva." Orig. art. has: 3 formulas and 1 figure.

Cord 2/K Inst of Chemical Phipins

KISSIN, YU.V.; HELOYE, G.P.; YEREHURA, I.V.; VELICEER ECVA, Ye.A.; TOVETECVA, V.I.; CHERKOV, N.M. Spectroscopic criterion of the isotacticity of polypropylene.

Vyskom.soed. 5 no.7:1117 Jl '63. (CIIA 16:9)

(Propylene—Spectra)

> CIA-RDP86-00513R000722910012-7" APPROVED FOR RELEASE: 09/17/2001

<u>1 1667-63</u>

EPR/ENP(j)/EPF(c)/EVT(m)/BDS AFFTC/ASD Ps-4/Pc-4/Pr-4

5/076/63/037/004/007/029

7.3

AUTHOR:

Khodzhayeva, I. V., Kissin, Yu. V.

7/

TITLE:

Effect of the structure of diethyldithiocartamates on the nature of

their isotopic exchange with \$35

PERIODICAL:

Zhurnel fizicheskoy khimii, V. 37, No. 4, 1963, 791-796

Dithiocarbanates are of interest because of their use in the rubber industry to accelerate the vulcanization of rubber and in agriculture as plant growth stimulators, insecvolungicides, and wood-decay inhibitors as well as reagents in organic chemistry. Properties of dithiocarbanates are discussed and the results are given of an investigation of the effect of the atructure of dithiocarbanates on the nature of their isotopic exchange with radioactive tetraethyl-thiurandiculfide. V Isotopic exchange of tetramethylthuriandisulfide and dithiocarbanates with S35 takes place at 120-180 degrees with an activation energy of 22-36 Kcal/mole. Tetramethylthuriandisulfide undergoes exchange with S35 more readily than the dithiocarbanates since the reaction is accompanied by the rupture of only the C-S bond, whereas in the exchange of dithiocarbanates with S35 the Me-S bond must be broken. The more polar the bond, the more it promotes exchange with S35. Steric hindrances play an essential part in the exchange reaction

Card 1/2

L. 16917-63

8/076/63/037/004/007/029

Effect of the structure of diethyldithiocarbamates on ...

2_

of Fe and Co dithiocarbamutes. Dithiocarbamates with a central atom having a coordination number of 4 exchange almost instantaneously with tetraethylthuriamdisulfide. If the coordination number is 6, the exchange is either much more difficult or does not take place at all. There are 2 tables. The authors express their gratitude to corresponding member of the Academy of Sciences USSR Ya. K. Syrkin for his assistance in their work.

ASSCCIATION: Moskovskiy institut tonkoy khimicheskoy tekhnologii imeni M. V.

Lomonosova (Moscow Institute of Fine Chemical Technology imeni

M. V. Lomonosov), Moscow

SUEMITTED: March 8, 1962

Card 2/2

KISSIN, Yu.V.; TSVETKOVA, V.I.; CHIRKOV, N.M.

Determination of the degree of isotacticity of polypropylene from its infrared spectra. Dokl. AN SSSR 152 no.5:1162-1165 0 '63. (MIRA 16:12)

1. Institut khimicheskoy fiziki AN SSSR. Predstavleno akademikom N.N.Semenevym.

8/0190/64/006/005/0962/0963

ACCESSION NR. AP4037293

AUTHORS: Zharov, A. A.; Kissin, Yu. V.; Pirogov, O. N.; Yenikolopyan, N. S.

TITLE: Radical stereospecific high pressure polymerisation of propylene

SOURCE: Vywsokomolekulyarnywye soyedineniya, v. 6, no. 5, 1964, 962-963

TOPIC TAGS: propylene polymerization, high pressure polymerization, radical stereospecific polymerination, isotactic propylene polymer

AISTRACT: Isotactic polypropylene was obtained by radical polymerization of propylene at 7000 atmospheres pressure and at temperatures of 100 or 200C. The polymerization of propylene occurs in the presence of such initiators as azobutyromitrile, benzoyl peroxide, and tert.butylperoxide (as well as without them). The molecular weight of the polymer obtained at 2000 in the presence of benzoyl peroxide was 900. Infrared spectroscopy showed that the polymer was in a state of isotactic configuration. This was confirmed by x-ray photographs. The polypropylene obtained by radical polymerization at 2000 was 15-19% isotactic, while the one obtained at 1000 was 54-56% isotactic. The degree of crystallinity

Card 1/2

CIA-RDP86-00513R000722910012-7" **APPROVED FOR RELEASE: 09/17/2001**

FIRSOV, A.P.; KISSIN, Yu.V.; CHIRKOV, N.M.

Stereospecificity of the W-TiCl₃ - Me(C_2H_5)n in the polymerization of propylene as dependent on the nature of metal of the metalloorganic compound. Vysokom.soed. 6 no.8:1537-1538 Ag 464 .

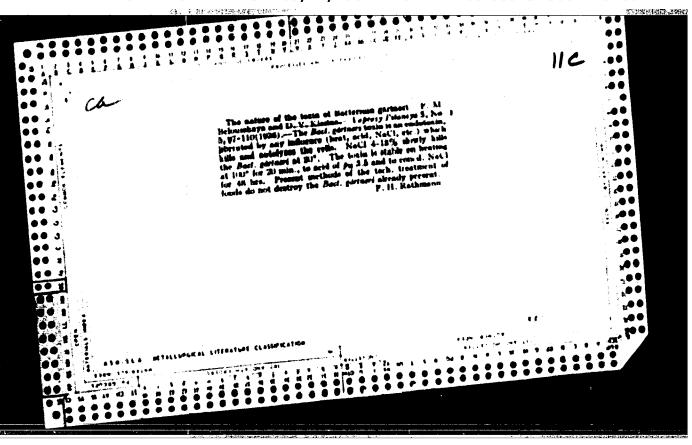
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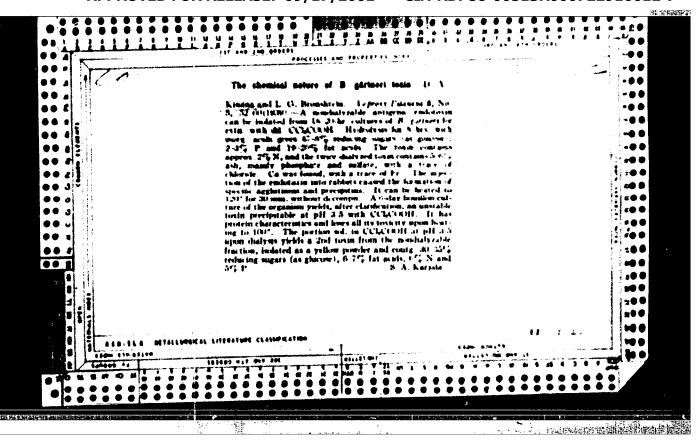
1. Institut khimicheskoy fiziki AN SSSR.

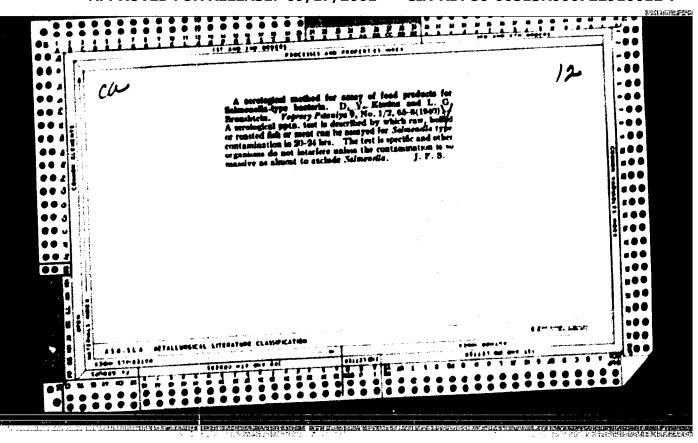
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KISSIN, Yu.V.: TOVETKOVA, V.I.; CHIRKOV, N.M. Determination of the isotacticity of polypropylene by means of infrared spectroscopy. Vysokom. sced. 7 no.71)288-2.90 J1 165. (MIRA 18:8) 1. Institut khimicheskoy fiziki AN SSSR.







"APPROVED FOR RELEASE: 09/17/2001

CIA-RDP86-00513R000722910012-7

TELEPHANE D. T.

USSP/Medicine - Fungi Medicine - Antiserum

Aug 48

"Studies of the Scrological Properties of the Fusarium Fungus, Isolated From Herbs Which Remain Through the Winter Under the Snow Cover," V. G. Geymberg, D. V. Lissina, Sector of Mutritional Hygiene, Inst of Mutrition, Acad Ked Sci USSE, 5 3/4 pp

"Gig i San" No C

Obtained antiserum through injections of extracts in rabbits. Explains use of the moldy growth of liquid culture of Fusarium Fungus in preparation of aqueous-saline extracts. Discloses reactions obtained. Includes four tables.

PA 28/1:9T00

Funri

"Certain properties of the "toxin' Fusarium shorotrichioides." Yu. I. Rubinsteyn. by D.V. Kissina. Gig. i san., No. 2, 1952.

Honthly List of Russian Accessions, Library of Congress, June 1952. UNCLASSIFIET.

"APPROVED FOR RELEASE: 09/17/2001 CIA-RDP

CIA-RDP86-00513R000722910012-7

Joint - Diseases

"Experimental alimentary mycotoxic endochondal osteodystrophia; on the etiology of Kaschin-Beck disease." M.I. Razumov, Yu.I. Rubinshteyn. Reviewed by D.V. Rissina. Gig. 1 san, No. 2, 1992.

Monthly List of Bussian Accessions, Library of Congress, June 1952, Unclassified

Kale

"STudy of the assimilability and of the food value of sea kale." A.F. Lerun,
O.P. Molchanova. Reviewed by D.V. Kissina. Gir. i san., No. 2, 1052.

Monthly List of Russian Accessions, Library of Congress, June 1752. UNGLASSIFIED.

KIRTIPHA, D.V.

Food- Preservation

Annotations. Gig. 1 san. no. 3, 1952

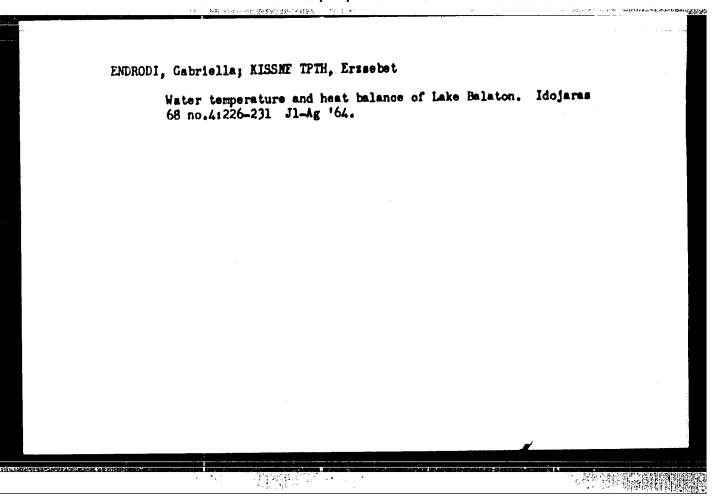
Monthly list of Bustian Accessions, Library of Congress, August 1952. INCLASSIFIED.

KISSINA, L.B.; TALOV, N.P.

Nature of the square mark of high etchability on 1Kh18N9T steel pipe blanks. Stal' 23 no. 3:263-266 Mr '64. (MIRA 17:5)

1. Zavod "Dneprospetsstalt" i TSentral'nyy nauchno-issledovatel'-skiy institut chernoy metallurgii imeni I.P.Bardina.

Device for automatic stopping of conveyers. Der. prom. 10 no.7:12-13 J1 '61. (MIRA 14:7) (Conveying machinery) (Automatic control)



KISSNE TOTH, E.

Climatological atlas of Africa. Idojaras 68 no.5:318
S-0 '64.

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

Features of the water cycle of dark soils of large sunken areas planted with trees. Trudy Inst. less 36:99-112 '56. (MRA 11:10) (Soil moisture) (Caspian Depression--Forest soils)

_118515, T.Ya.

Results obtained in observing the effect of forest masses and strips on the distribution of snow and the absorption of moisture by soil in spring. Trudy Inst.less 43:138-151 158.

(MIRA 11:12)

(Forest influences) (Soil moisture) (Snow)

APPROYED: EQB: HELEASE: PQ9/17/2001ctur.GIA-R-DR86; AQ5,13R000722910012-7

Abs Jour: Ref Zhur - Fizika, No 7, 1958, No 15552

Author : Kinsler Fordinand M.

Inst : Ustav po vyzkum a vyuziti paliv, Prague

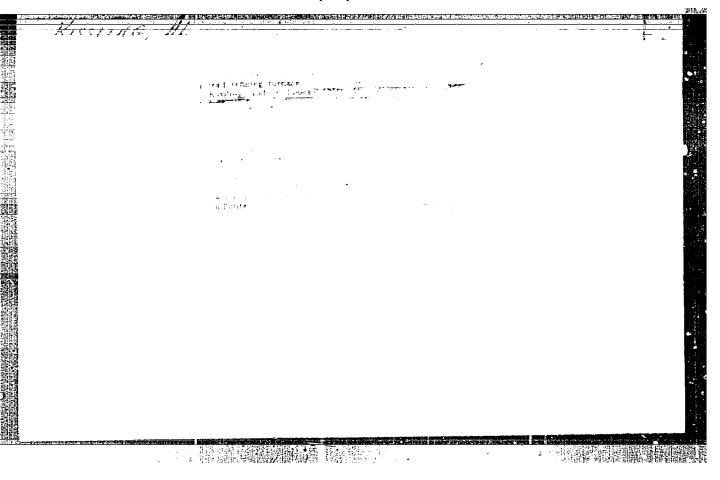
Title : Determination of the Changes in the Lattice Parameter of

Various Graphitas.

Orig Pub: Chem. listy, 1957, 51, No 1, 13-20

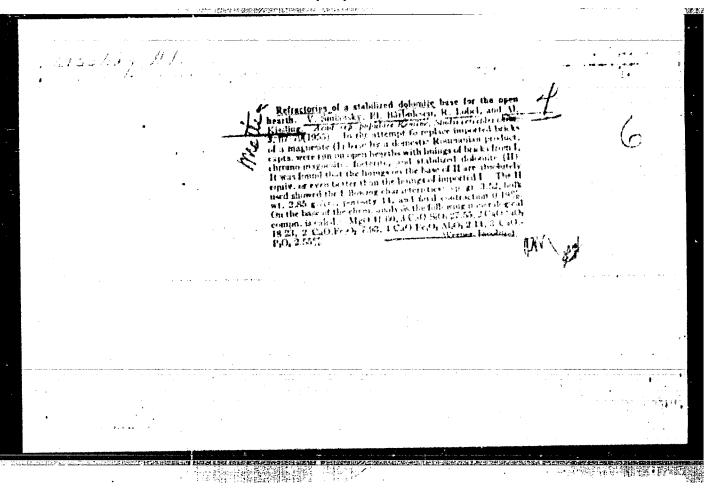
Abstract: Description of a new method for determining the lattice parameters of graphite, using an asymptrical base for the photographic film and radiation with a relatively large wavelength (\(\frac{1}{2} \) Cr). It is indicated that the method of backward reflection does not justify itself in the study of graphite. The accuracy of determining the interplanar distances by the proposed method is \(\frac{1}{2} \) 0.003 \(\hat{L} \). The method gives very exact results in the study of the changes of the interplanar distances of carbons in graphitization and in the detection of the rhombohedric structure of graphite.

Card : 1/1



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KISSLING,

RUHANIA / Cosmochemistry. Geochemistry.

Hydrochemistry.

: Referat Zhur--Khimiya, No. 11, 1959, 38157 Abs Jour

: Codarces, A.; Kissling, A.; and Kissling, M. : Rumanian Academy of Sciences Author

: Note on the Ludwigite from Ocna de Fiers Inst

Title

: Bull Stiint Acad RPR, Sec Geol Si Geograph, 2, No. 3-4, 515-527 (1957) (in Rumanian with summaries in French and Russian) Orig Pub

: Using the microscopic and especially the chemical method, the authors have studied Ludwigite asso-Abstract

clated with magnetite in a skarn formation in which the authors have also identified serpentine, fosterite, ascharite, hematite, pyrite, sphalerite, chalcopyrite, quartz, and limonite. The chemical composition of the dolomite is as

Card 1/3

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RULLANIA / Cosmochemistry. Geochemistry.

Hydrochemistry.

: Referat Zhur--Khimiya, No: 11, 1959, 38157 Abs Jour

: Codarcea, A.; Kissling, A.; and Kissling, M. : Rumanian Academy of Sciences : Note on the Ludwigite from Ocna de Fier. Author

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SCCCIESCU, M.; DIACONU, F1.; KISSLING, M.

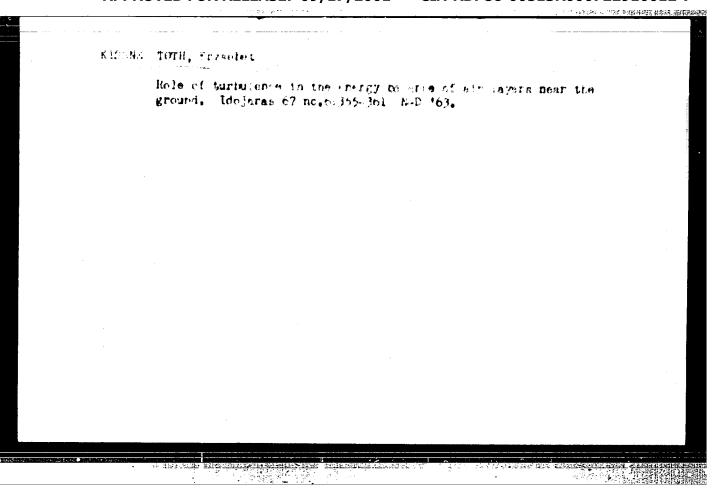
Contributions to the knowledge of the genesis of the mineralization in the Blasma Valley. Rev min 12 no.6:253-258 Je '61.

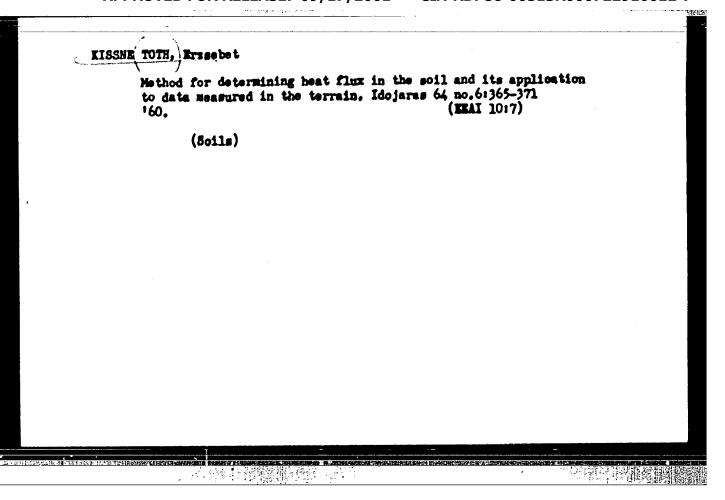
APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

KISSNE TOTH, Erzsobet

Role of turbulence in the formation of energy balance in the air layers situated above the surface of lakes. Orsz meteor int besz tud kut 26:284-290 162(publ. 163).

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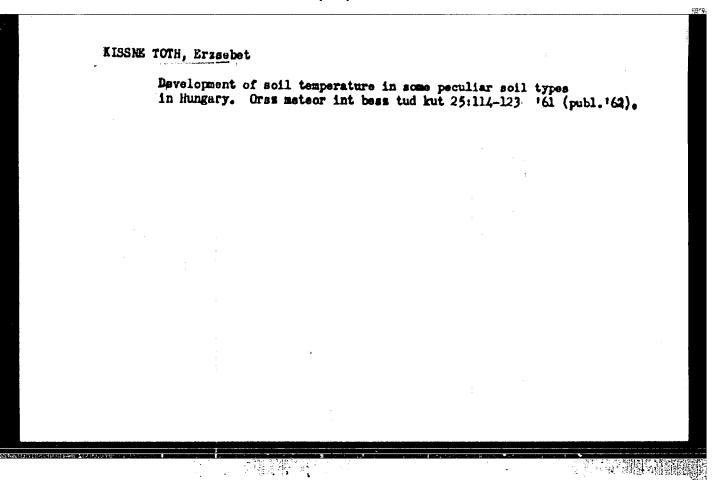


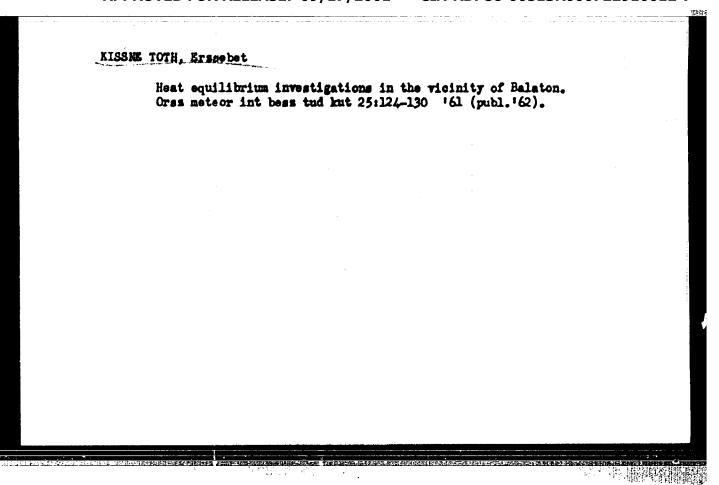
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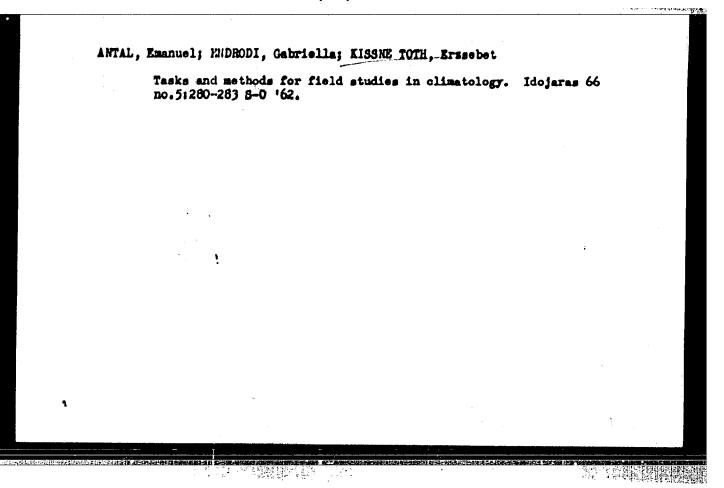
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KISSI : TOTH, Errsebet

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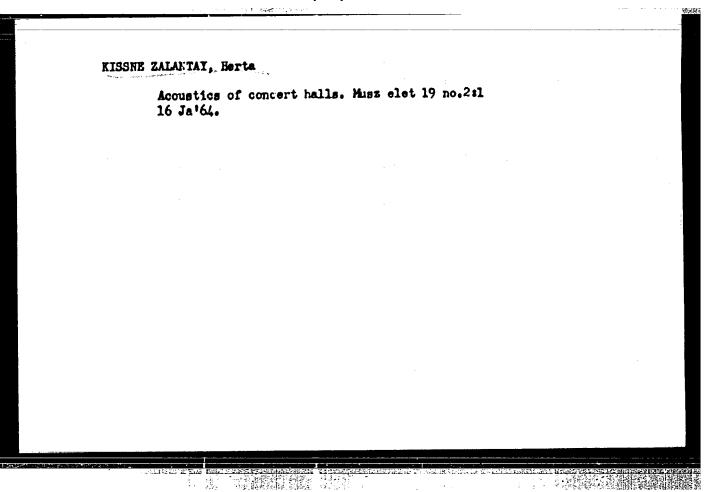
FAREDIN, Imre; KISSNE SZABADAI, Iren; WINTERNE SIMOR, Ilona technikai segedletevel

Simple method for the determination of the 17,21-dihydraxy-20-ketosteroid content of the urine by means of the Porter-Silber color reaction. Kiserl. cevostud. 14 no.5:549-555 0 '62.

1. Szegedi Orvostudomanyi Egyetem I. sz. Belgyogyaszati Klinikaja. (HYDOCORTISONE) (URINE)

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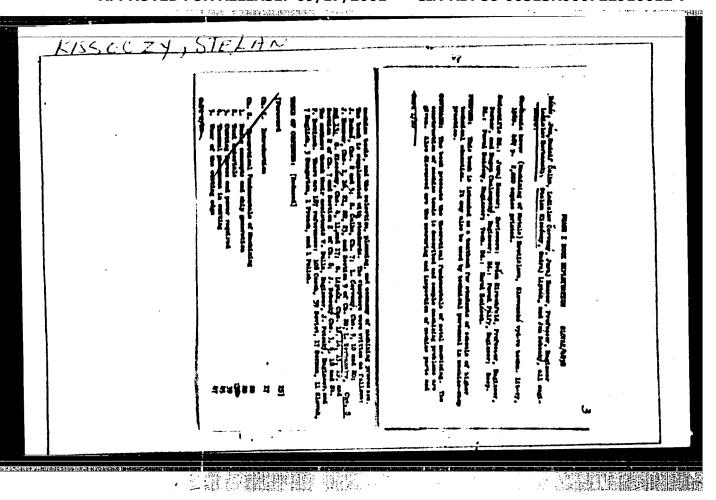
Improving the productivity in grinding. p. 526. (TECTOTICKA PRACA, Vol. 9. No. 3, Aug 1957, Bratislava, Gzechoslovakia)

CO: Konthly List of East Suropean Accessions (SEAL) LC. Vol. 6, No. 12, Dec 1957. Uncl.

A contribution to chip forming in turning operations. r. 535.

(TECHNICKA PRACA, Vol. 9, No. 8, Aug 1957, Bratishava, Vzechoslavakia)

50: Yonthly List of Past European Accessions (LEAL) i.C., Vol. 6, No. 12, Wee 1957. Uncl.



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KISSOCZY, S.

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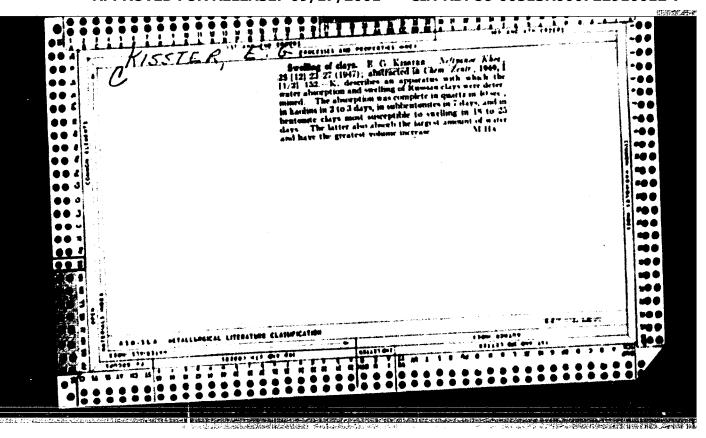
CZECHONLOYAKIA

KISSOCZY, S; STERBOVA, S.

1. Internal Medicine Department of the Okres Institute of National Health (Interne oddelenie Okresneho ustavu narodneho zdravia), Presov (for Kossoczy);
2. Internal Medicine Chair SUDL (Interna katedra SUDL), Trencin

Bratislava, Lekarsky Obzor, No 2, 1963, pp 65-69

"Confidence in the Physician."



GENIYEV, G.A., doktor tekhn.nauk; KISSYUK, V.N., insh. A basis for the conditions affecting concrete strength.

Bet. i zhel.-bet. 8 no.12:553-557 D 162. (MIRA 16:2) (Concrete-Testing)

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

BRUTYO, Janos; TENYI, Ferenc, technologus; MARTIN, Jancs; KIS SZABO, Laszlone; ARADI, Tibor; HOFFMANN, Nandor; KIRALY, Albert; BOROSS, Istvan, mernok

National conference of socialist brigade leaders. Munka 15 no.4: 10-17 Ap '65.

1. Secretary General of the Contral Council of Hungarian Trade Unions, Budapest (for Brutyo). 2. Lang Machine Factory, Budapest (for Tenyi). 3. Tatabanya Coal Mining Trust, Tatabanya (for Aradi). 4. Kobanya Drug Factory, Budapest (for Hoffmann). 5. Research Institute of Heavy Chemical Industry (for Kiraly). 6. Csepel Automobile Factory, Budapest (for Boross).

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

KISSZEKELYI, Odon, Dr.; THENCSENI, Tibor, Dr.

Case of meningoencephalitis detected in the acute phase and caused by Gryptococcus neoformans in Hungary. Orv. hetil. 98 no.40: 1110-1112 6 Oct 57.

1. A Magyar Mephadsereg Egessegugyi Ssolgalatanak koslemenye.

(MENINGOENGEPHALITIS, etiol. & pathogen.

Cryptococcus neoformans, histopathol. (Hun))

(CHYPTOCOCCOSIS, case reports

meningoencephalitis caused by Cryptococcus
neoformans, histopathol. (Hun))

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

\$/166/62/000/001/003/009

AUTHORS:

Kist, A. A., Lobanov, Ye. M., Zvyagin, V. I., Bartnitskiy,

TITLE:

Effect of gamma irradiation upon oxide films of germanium PERIODICAL: Akademiya nauk Uzbekskoy SSR. Izvestiya.

matematicheskikh nauk, no. 1, 1962, 88-90

TEXT: The effect of gamma rays on germanium monoxide and germanium Seriya fizikodioxide films produced by etching was quantitatively measured with a Geirovskiy micropolarograph. The monoxide - dioxide mixture produced by etching germanium powder in standard etching agent did not change under gamma irradiation in air, carbon dioxide, and in vacuum (10-4 torr) with 20, 60, 100, 150, and 200 million r. In the subsequent irradiation of the weighed portion of germanium etched in a standard reagent with 20, 30, 50, and 100 million r, the amount of germanium dioxide increases at doses of up to 40-50 million r, and then decreases again. The oxide film produced in etching agent no. 5 contains monoxide and dioxide in and : 1 ratio. While etching agent no. 5 gives rise to germanium monoxide,

3/166/62/000/001/009/009 B125/B104

Effect of gamma irradiation ...

germanium dioxide is contained in the film in an equal amount. The anomalous current and the photocurrent are not exclusively due to the germanium monoxide. Similar phenomena are also observed when exposing the diodes to gamma irradiation (doses above 106 r). These anomalies disappear either entirely or partially at doses of more than 108 r. The irradiated photodiodes yield a photocurrent at such doses if the amount of germanium dioxide on the surface increases. The upper limit of the anomalous photocurrent shifts toward the visible region when etching agent no. 5 is used. Gamma irradiation first causes the oxide film to grow more considerably, but the secondary fast electrons then again partly destroy the oxide film. As a result, the oxide film becomes eventually thinner. If present considerations are correct, germanium diodes are made insensitive also to intense radiations in that the oxide film is prevented from growing all throughout the dose range. There are 1 figure, 1 table, and 8 references: 2 Soviet and 6 non-Soviet. The four references to English-language publications read as follows: S. I. Ellis, Appl. Phys. 1957, 11, 1262, 28; I. Everest, J. Chem. Soc., Febr. 1953, 660; I. Bardet, Tchakarian A. C. R., 1928, 637, 186; L. Dennis, Xules R. J. Am. Soc., 1930, 3554, 52.

Card 2/3

Effect of gamma irradiation ...

S/166/62/000/001/009/009 B125/B104

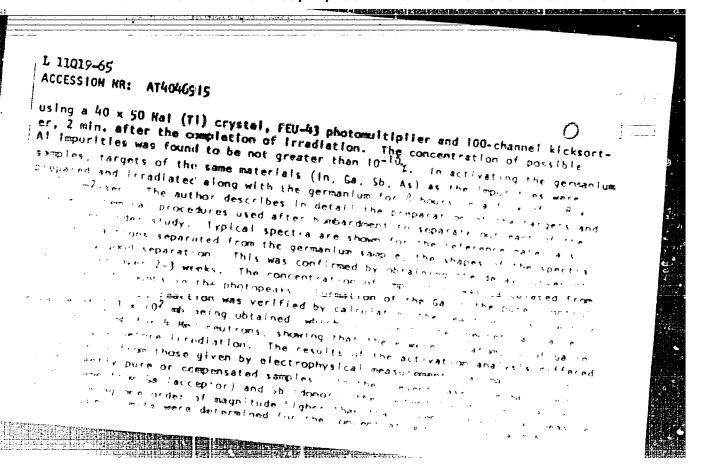
ASSCCIATION: Akademiya nauk UsSSR (Academy of Sciences of the

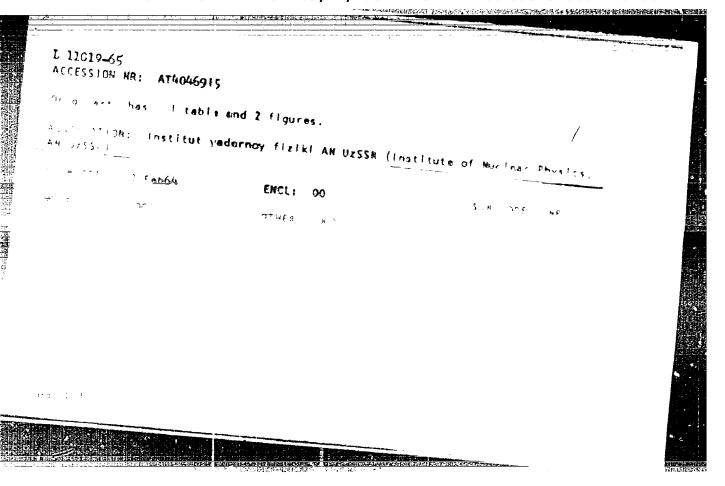
Uzbekskaya SSR)

SUBMITTED: August 25, 1961

Card 3/3

ACCESSION NR: AT4046915 S/0000/64/000/000/000/000/000/0001/0001 AUTHOR: Lobanov, Va. M.: Ivyagin, V. I.: Kist, A. A.: Sviridova, A. I.: Vevseyenko, M. I.: Vevseyenko, J. I.: Kist, A. A.: Sviridova, A. I.: Vevseyenko, J. I.: Kist, A. A.: Sviridova, A. I.: Vevseyenko, J. I.: Kist, A. A.: Sviridova, A. I.: Vevseyenko, J. I.: Kist, A. A.: Sviridova, A. I.: Vevseyenko, J. I.





DULOVA, V.I.; KIST, A.A.; LEONT'IEV, V.B.

Interaction of ions and molecules of some acids with cyclohexanol.

Interaction of ions and molecules of some acids with cyclohexanol.

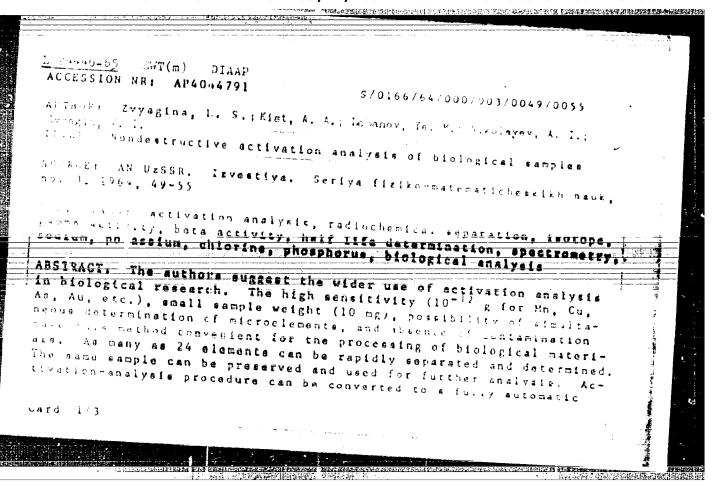
Interaction of ions and molecules of some acids with cyclohexanol.

(Acids, State of the cyclohexanol)

(MIRA 15:12)

Kafedra hoorganicheskoy khimii.

(Acids, Organic) (Cyclohexanol)

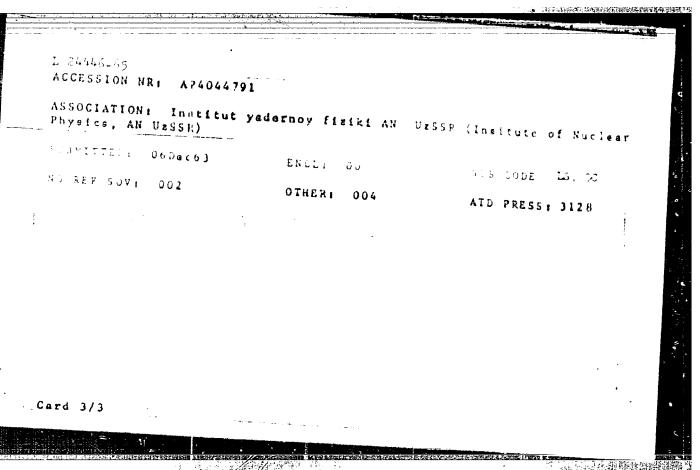


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CIA-RDP86-00513R000722910012-7

ACCESSION NR: AP4044791 system. Automatic units for irradiation, activity counting, and dat processing have already been developed. The basic problem in this analysis is the deparation of the activity of a given elegent. This problem can be solved by chemical separation, invatification from . spectra, B activ ty, or half life, etc., or colons the temple. The authors used activate a writer to a field the rate of healthy and cancerous rate, littlediation min in a neutron flux (1.8 x 10 cm⁻² sec⁻¹ and 1.2 x 10 cm⁻² cml'secl'), for determination of sodium, chlorine, potassium, and phosphorus. Activities of these elements were measured by means of a verpectrometer, a Beanalyzer constation of the recent city and x limi, an FTI-11 photomultiplier, and a ru-, by sudiameter. A desett, from is given of the method used. The account with the determiration fails in the 5-102 error range (e.g.,) and life for K" was 11.3 mil. o hr, as compared to 12.5 hr). The number of elements determined in nondestructive analysis can be ausmedied by the removal y. Wa . rom the semple after irradiation, and by the use of anticoincitations, for, and B-y coincidence scheme; developed for this purpose, magnetic analyzers, resonance irradiation, etc. Orig. art. has: 5 Card 2/3; TO CARESTOLING CONTRACTOR OF THE PARTY OF TH

"APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7



LOBANOV, Ye.M.; ZVYAGIN, V.I.; KIST, A.A.; ZVEREV, B.P.; SVIRIDOVA, A.I.; MOSKOVTSEVA, G.A.

Determination of manganese in silicon by the radioactivation method. Zhur. anal. khim. 18 no.11:1349-1355 N '63.

1. Institut yadernoy fiziki AN UzSSR, Tashkent. (MIRA 17:1)

KIST, A.A.; AVYAGINA, L.S.; LOBANOV, Ye.M.; GVIRIDOVA, A.I.: MOSKOVISEVA, G.

Activation analysis of copper and mangamene in biological objects. Izv. AN Uz. SSR. Ser. fiz.-mat. nauk 8 no.2:77-80 '64. (MIRA 17:9)

1. Institut yadernoy fiziki AN UZSSR.

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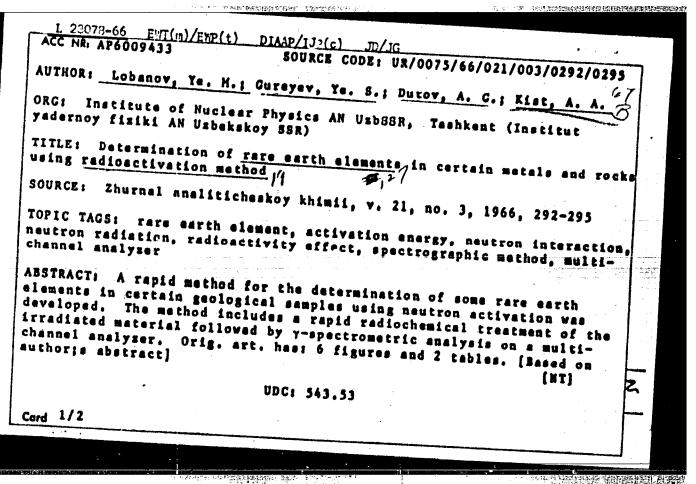
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MIST, A.A.; ZVYAGINA, L.S.; LOBANOV, Ye.M.; MOSKOVTSEVA, C.A.

Determination of halogens in biological materials by the activation method. Zhur. anal. khim. 20 no.11112-117 '65. (MIRA 18:3)

1. Institut ymdernoy fiziki AN UzSSR, Tashkent.



"APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7

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ACC NR: AP7008895

SOURCE CODE: UR/0425/66/009/009/0012/0016

AUTHOR: Lobanov, Ye. M.; Khotamov, Sh.; Kist, A. A.

CRG: Physics-Engineering Institute im. S. U. Umnrov, AN TadzhSSR (Fiziko-tekhnicheskiy institut AN TadzhS3R); Nuclear Physics Institute, AN UzSSR (Institut yadernoy viziki

TITLE: Determination of certain rare-earth elements in the ash of plants and soils

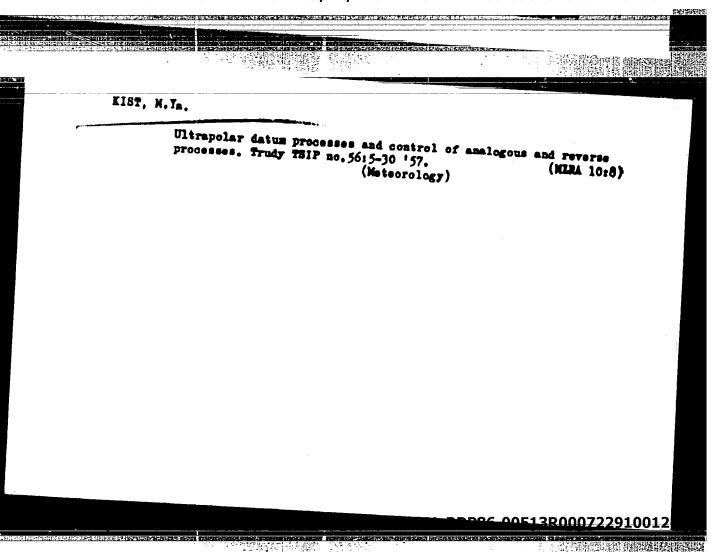
SOURCE: AN TadzhSSR. Doklady, v. 9, no. 9, 1966, 12-16

TOPIC TAGS: gamma spectrum, neutron irradiation, rare earth element, radioisotope,

SUB CODE: 06, 18, 20

ABSTRACT: Radiation of Artemisia terrae albae wormwood ash in a stream of 1.8x10¹³ neutrons/cm², with 40-hour holding period, is sufficient for determination. Prolonged "cooling" prevents determination of short-lived isotopes. A complete 7 -ray spectrum of the sample was used and decay curves were plotted for accurate identification of individual Y emitters and separation of individual photopeaks, followed by graphical analysis. From the Compton distribution of N_a^{24} and Sc⁴⁰ the contribution from N_a^{24} was determined by comparison with a standard. Results were compared with those from radiochemical separation of A. A. Adkhamov, Corresponding Member, Tadzhik Academy of Sciences, 19 March 1966. · Card 1/1 UDC: none

MARKA MARKA MARKA



10FFE, Ya.A.,; NIKONOVA, I.I.; CHERTKO, V.F.; HAYDENOV, G.N.; ZIKIN, B.N.; NOCHEVKINA, L.P.; NESTEROV, L.I.; KISTANOV, H.I.; KUDROV, V.M.; FAK, G.V., red.; PONOMAREVA, A.A., tekim. red.

[Structural changes in the industries of the United States, Great Britain and German Federal Republic in the postwar year]Strukturnye izmoneniia v promyshlennosti SShA, Anglii i FRR v poslevoennye gody. Monkva, Ekonomizdat, 1962. 417 p.

[Mina 15:10]

[United States—Industries] (Great Britain—Industries)

(Germany, West—Industries)

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"APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7

Corn in Thunns. Zemladelie 26 no.3:41-43 Mr U.A.

1. Malayakaya opytno-meliorativnaya stantsiya imeni

prof. r.A. Kontychova.

Relation of rice to the salinity of soils. Pechvovedenie no.5:

(#IRA 16:5)

1. Vallyskaya opytno-meliorativnaya stantsiya imeni

P.A.Kostycheva.

(Volga-Akhtuba floodplain—Rice)

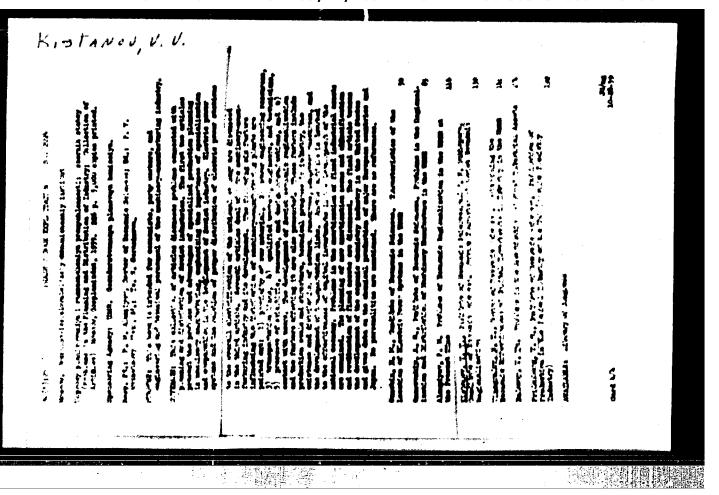
(Volga-Akhtuba floodplain—Saline and alkali sails)

CHERNYKH, A.A., kand.tekhn.nauk; KISTANOV, N.S., kand.tekhn.nauk
Recharging natural limans. Gidr. i mel. 16 no.1:12-17 Ja '64.

(MIRA 17:2)

APPROVED FOR RELEASE: 09/17/2001 CIA-RDP86-00513R000722910012-7"

KISTANOV, V.V.; KRIVTSOV, S.G.; SPIDCHENKO, K.I.; SUKHOPARA, F.W. "Roonomic geography of the Soviet Union: Russian Soviet Federative Socialist Republic." Reviewed by V.V. Kistanov and others. Inv. AN 88SR. Ser. geog. no.4:128-132 J1-Ag 157. (MIRA 11:1) (Geography, Economic) (MIRA 11:1)



3(5)

SOY/10-59-3-7/32

AUTHOR:

Kistanov, V.V.

TITLE:

Some Particularities of the Formation of the Economic Areas

in the East of the Country

PERIODICAL:

Izvestiya Akademii nauk SSSR, Seriya geograficheskaya, 1959,

Nr 3, pp 62-67 (USSR)

ABSTRACT:

Forty percent of all Soviet investments in 1959-65 will be used in the Eastern areas. For the time being, although the East areas (Urals and Soviet Asia) cover about 4/5 of the total Soviet territory and possess 3/4 of the country's thermo-power and raw material resources, they only have 1/3 of the population and 1/4 of its industry. Yet, the increase of heavy industry in those regions is The Urals region has now (1957) 82 times more heavy industry than it had in 1913. West Siberia has 204 times more, Kazakhstan 97 times more; East Siberia 58 times more, Far East 50 times more. The author distinguishes 3 groups of Eastern areas, each group consisting of 2 large economic-geographic areas: 1) Urals and West Siberia; 2) Kazakhstan and Central

Card 1/4

507/19-59-3-7/32

Some Particularities of the Formation of the Economic Areas in the East of the Country

> Asia; 3) East Siberia and the Far East. Each of these areas is described from the standpoint of the national economy. The statistical bureau of TsSU USSR has stated that of the 29 branches of the machine building industry, the following number of branches have been installed in the areas, (Urals - 26; West Siberia - 25; Kazakhstan - 17; Central Asia - 21; East Siberia - 13; Far East - 12). By 1965, the following sources of energy will be put into operation in the East: the Bratckaya GES (3.6 million kW); Nazarovskaya GRES (1.2 million kW). The powerful Krasnoyarsk GES is under construction; and the Irsha, Itat and other thermoelectric plants will be built. The electric capacity of the East-Siberian area will be increased by almost 7 times by 1965. This energy will be very cheap (1 kW/h for 1.5 kopecks). Central Siberia will have its own huge power system (the power plants on the Angara and the Yenisey rivers) utilizing about 75 to 80% of the capacity of the area. Almost all of the Eastern areas will have their own powerful non-ferrous metallurgy (espe-

Card 2/4

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Some Particularities of the Formation of the Economic Areas in the East of the Country

cially the Angara-Yenisey area with its aluminum, magnesium and titanim, and Kazakhstan). The 3rd ferrous-metallurgical base of the country will be built in Western Siberia. . Kazakhstan, the Angara valley and the Trans-Baykal region will have particular importance because of their iron resources and plants. The Karaganda and the Tayshet plants are mentioned by names. Several paragraphs are devoted to the importance of the communications network in the East. Mentioned is the Lena RR which made the construction of the Bratskaya GES, the Korshunovskiy gornoobogatitel'nyy kombinat (Korshinovskiy Mining and Ore-Concentrating Combine), a large wood-processing center, and the organization of the entire Bratskaya-Tayshet industrial area possible. The Achinsk-Abalakovo RR, now under construction, will be lengthened, so as to reach the region beyond Angara. The Hizhne-Angarskiy (Lower-Angara) industrial center is under construction. The Bam-Chultman RR, now under construction, will later be lengthened to Yakutsk and Magadan. In this way, the format-

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ion of the North-Eastern economic area will be stepped up. The author states that contemporary current attempts to divide the Soviet East into large economic areas are lacking in sufficient knowledge of the sites and their industrial possibilities. There are 4 Soviet references.

ASSOCIATION:

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